

Medical Diagnosis/Conditions for Eligibility in AEIS

- 1) Achondroplasia
- 2) Agenesis of Corpus Callosum
- 3) Agyria (Lissencephaly)
- 4) Albinism
- 5) Amniotic Band syndrome
- 6) Anencephaly
- 7) Angelman's syndrome
- 8) Anophthalmia
- 9) Apert syndrome
- 10) Aplasia of the brain (brain malformation/abnormality)
- 11) Arhinencephaly (Holoprosencephaly)
- 12) Arnold-Chiari syndrome
- 13) Arthrogyposis
- 14) Asperger syndrome/disorder
- 15) Asphyxiating Thoracic Dystrophy (Jeune syndrome)
- 16) Attachment disorder
- 17) Autism/Autism Spectrum disorder
- 18) Bardet-Biedl syndrome
- 19) Brain injury/degeneration
- 20) Brain malformation/abnormality
- 21) Cerebral Palsy (all types)
- 22) CHARGE syndrome
- 23) Chiari Malformation
- 24) Childhood Depression
- 25) Childhood Disintegrative disorder
- 26) Cornelia de Lange syndrome
- 27) Cortical vision impairment (vision loss/impairment)
- 28) Cri-du-Chat syndrome
- 29) Cytomegalovirus (CMV)
- 30) Dandy Walker syndrome/variant
- 31) De Morsier syndrome (Septo-Optic Dysplasia)
- 32) Developmental Apraxia
- 33) DiGeorge syndrome
- 34) Dilantin syndrome (Fetal Hydantoin syndrome)
- 35) Down Syndrome (Trisomy 21)
- 36) Edwards syndrome (Trisomy 18)
- 37) Encephalomalacia
- 38) Encephalopathy
- 39) Epilepsy (seizure disorder)
- 40) Fetal Alcohol syndrome
- 41) Fetal Hydantoin syndrome (Dilantin syndrome)
- 42) Fragile X syndrome
- 43) Genetic/Chromosomal malformation/abnormality (not listed)
- 44) Hearing Loss/Impairment
- 45) Heart Disease/Defect (not listed)
- 46) Hemiplegia
- 47) Herpes Simplex Virus (HSV)
- 48) Holoprosencephaly (Arhinencephaly)
- 49) Holt Oram syndrome
- 50) Hydranencephaly
- 51) Hydrocephaly (with or without shunt; congenital or acquired)
- 52) Hypoplastic Left Heart syndrome
- 53) Incontinentia Pigmenti
- 54) Infantile spasms (seizure disorder)
- 55) Intraventricular Hemorrhage (IVH) Grade III or IV
- 56) Jeune syndrome (Asphyxiating Thoracic Dystrophy)
- 57) Klinefelter's syndrome
- 58) Landau-Kleffner syndrome
- 59) Lead (elevated blood levels)
- 60) Leber Congenital Amaurosis (vision loss/impairment)
- 61) Leukodystrophy
- 62) Lissencephaly/Lissencephaly syndrome (Agyria/Miller-Dieker syndrome)
- 63) Macroencephaly (brain malformation/abnormality)
- 64) Macrogyria (brain malformation/abnormality)
- 65) Megalencephaly (brain malformation/abnormality)
- 66) Meningitis
- 67) Meningomyelocele (Spina Bifida)
- 68) Menkes syndrome
- 69) Mercury Poisoning

- | | |
|---|--|
| 70) Microcephaly (brain malformation/abnormality) | 104) Seizure disorder/uncontrolled or poorly controlled seizures |
| 71) Microgyria (brain malformation/abnormality) | 105) Septo-Optic dysplasia (De Morsier syndrome) |
| 72) Mobius syndrome | 106) Shaken Baby syndrome |
| 73) Mucopolysaccharidosis | 107) Smith-Lemli-Opitz syndrome |
| 74) Muscular Dystrophy | 108) Spina Bifida (Myelomeningocele) |
| 75) Myasthenia syndrome-congenital | 109) Spinal Muscular Atrophy (SMA) |
| 76) Myelomeningocele (Spina Bifida) | 110) Spinocerebellar Ataxia |
| 77) Myopathy | 111) Stickler syndrome |
| 78) Neurological Disease/Defect (not listed) | 112) Stroke, Prenatal or Neonatal |
| 79) Neurofibromatosis | 113) Sturge Webber syndrome |
| 80) Noonan's syndrome | 114) Tay-Sachs Disease |
| 81) Opitz G/BBB syndrome | 115) Traumatic Brain Injury (TBI) |
| 82) Optic nerve hypoplasia or atrophy | 116) Traumatic Retinal Detachment |
| 83) Osteogenesis Imperfecta | 117) Treacher-Collins syndrome |
| 84) Other (not listed) | 118) Trisomy 13 (Patau syndrome) |
| 85) Pallister-Killian syndrome | 119) Trisomy 18 (Edwards syndrome) |
| 86) Patau syndrome (Trisomy 13) | 120) Trisomy 21 (Down syndrome) |
| 87) Pediatric Human Immunodeficiency Virus (HIV)/Acquired Immune Deficiency Syndrome (AIDS) | 121) Tuberous Sclerosis |
| 88) Periventricular Leukomalacia (PVL) | 122) Turner syndrome |
| 89) Pervasive Developmental disorder (PDD and PDD-NOS) | 123) Ventriculomegaly |
| 90) Phelan McDermis syndrome | 124) Vision Loss/Impairment |
| 91) Phiffer syndrome | 125) Weaver syndrome |
| 92) Porencephalic Cyst | 126) Williams syndrome |
| 93) Prader-Willi syndrome | |
| 94) Prematurity (26 weeks or less gestation and/or 1000 grams or less and 18 months chronological age or younger) | |
| 95) Pseudothalidomide syndrome (Roberts syndrome) | |
| 96) Reactive Attachment disorder (RAD) | |
| 97) Retinoblastoma | |
| 98) Retinopathy of Prematurity (ROP) | |
| 99) Rett syndrome | |
| 100) Roberts syndrome (Pseudothalidomide syndrome) | |
| 101) Rubella-congenital | |
| 102) Rubenstein-Taybi syndrome | |
| 103) Schizencephaly | |